In the Claims

Please cancel claims 1- 47 and 65-66 prior to calculating fees. Please amend claims 50 and 64.

Please re-write the claims as shown below.

- 1-47. (Cancelled)
- 48. (Original) A sustained release device comprising an MPL inhibitory agent that reduces platelet count in a subject, wherein the agent is released for at least 7 days.
- 49. (Original) The sustained release device of claim 48, further comprising a blood modifying agent.
- 50. (Currently Amended) The sustained release device of claim 49, wherein the blood modifying agent is selected from the group consisting of an anti-coagulant agent, a fibrinolytic agent and an inhibitor of platelet function.
- 51. (Original) The sustained release device of claim 48, wherein the MPL inhibitory agent is released in an amount effective to reduce platelet count in a subject to below normal levels.
- 52. (Original) The sustained release device of claim 48, wherein the MPL inhibitory agent is released at a rate ranging from 0.01 μ g/kg/day to 30 mg/kg/day.
- 53. (Original) The sustained release device of claim 48, wherein the MPL inhibitory agent is released for at least 30 days.
- 54. (Original) The sustained release device of claim 48, wherein the MPL inhibitory agent is released for at least 6 months.
- 55. (Original) The sustained release device of claim 48, wherein the MPL inhibitory agent is released for at least 1 year.
- 56. (Original) The sustained release device of claim 48, wherein the MPL inhibitory agent is released for at least 5 years.
- 57. (Original) The sustained release device of claim 48, wherein the MPL inhibitory agent is released in an effective amount that does not affect platelet function.
 - 58. (Original) A pharmaceutical preparation comprising

an amount of an agent that inhibits signal transduction from an MPL receptor effective to reduce platelet count, and

a pharmaceutically acceptable carrier.

59. (Original) A pharmaceutical preparation comprising
an amount of an agent that binds to an MPL receptor effective to reduce
platelet count, and

a pharmaceutically acceptable carrier.

- 60. (Original) The pharmaceutical preparation of claim 59, wherein the agent binds to an extracellular region of an MPL receptor.
- 61. (Original) A pharmaceutical preparation comprising
 an amount of an agent that binds to a thrombopoietin molecule effective to
 reduce platelet count, and
 a pharmaceutically acceptable carrier.
- 62. (Original) A pharmaceutical preparation comprising
 an amount of an agent that binds to an intracellular tyrosine kinase that
 modulates signal transduction from an MPL receptor effective to reduce platelet count, and
 a pharmaceutically acceptable carrier.
- 63. (Original) A pharmaceutical preparation comprising
 an amount of an agent that inhibits binding of a thrombopoietin molecule to an
 MPL receptor effective to reduce platelet count, and
 a pharmaceutically acceptable carrier.
- 64. (Currently Amended) The pharmaceutical composition of claim 58, 59, 60, 61, 62 or 63, wherein platelet count is reduced to below normal levels.
 - 65.-66. (Cancelled)
- 67. (Original) A method for treating a subject having above normal platelet count comprising

administering to the subject in need of such treatment an MPL pathway inhibitory agent in an amount effective to reduce platelet count.